



Press release

Publication of the securities note having obtained visa no. 17-297 of the AMF in connection with the capital increase of GenSight Biologics

Paris, June 23, 2017 –GenSight Biologics (Euronext: SIGHT, ISIN: FR0013183985, PEA-PME eligible), (“GenSight Biologics” or the “Company”) informs that the listing prospectus containing the 2016 registration document of the Company, registered with the *Autorité des marchés financiers* (the “AMF”) on April 28, 2017 under number R.17-03, as well as the securities note containing a summary of the listing prospectus having obtained the visa n°17-297 on June 23, 2017 are made available to the public within the conditions of French regulations. A press release regarding the offering’s results was today published.

The funds raised will be allocated to prepare the launch of GS010 in Europe and the United States, and especially the financing related to the marketing and market access, as well as the establishment of a marketing infrastructure.

This capital increase reserved to a category of persons, of an amount of EUR 22,5 million representing approximately 19% of the share capital of the Company (on a non-diluted basis before the capital increase) benefited from the support of existing principal shareholders of the Company up to 33% of the offering in the following proportions:

<i>Shareholders > 5% of the share capital as of the launch of the offering and who participated to the capital increase</i>	Number of shares		
	Before Offering	After Offering	Subscription
Versant (via Venture Capital IV & Side Fund IV)	2,947,048	3,280,381	333,333
Abingworth Bioventures VI	2,873,306	3,139,973	266,667
Bpifrance Participations	1,500,000	2,000,000	500,000
Vitavest S.à.r.l	1,206,373	1,339,706	133,333
Total			1,233,333
<i>Other shareholders</i>			
Employees, Officers, Members of the Board of Directors	417,362	417,362	-
Others	10,840,164	13,356,831	2,516,667
Total	19,784,253	23,534,253	3,750,000

The shares were issued today by a decision of the Company’s Chief Executive Officer, pursuant to the authorization of the Board of Directors dated June 22, 2017 and the 22nd resolution of the extraordinary general meeting of the shareholders of the Company held on May 31, 2017 and in accordance with articles L. 225-138 of the French Commercial code (code de commerce). The capital increase was reserved to the categories of investors satisfying specific requirements as defined in the abovementioned 22nd resolution.

Risk factors

Attention is drawn to the risk factors related to the capital increase presented in Section 2 of the Securities Note and to risk factors related to the Company and its activities presented in Chapter 4 of its 2016 registration document. These documents are available free of charge from the Company website (www.gensight-biologics.com) and/or website of the Autorité des marchés financiers (www.amf-france.org).

This press release does not constitute a prospectus within the meaning of the Prospectus Directive or a public offering.

Contacts

GenSight Biologics

Thomas Gidoïn
Chief Financial Officer

tgidoïn@gensight-biologics.com

+33 (0) 6 01 36 35 43

RooneyPartners

Media Relations
Marion Janic

mjanic@rooneyco.com

+1-212-223-4017

The Trout Group

Investor Relations
Chad Rubin

crubin@troutgroup.com

+1-646-378-2947

About GenSight Biologics

GenSight Biologics S.A. (GenSight Biologics) is a clinical-stage biotechnology company discovering and developing novel therapies for neurodegenerative retinal diseases and diseases of the central nervous system. GenSight Biologics' pipeline leverages two core technology platforms, the Mitochondrial Targeting Sequence (MTS) and optogenetics for retinitis pigmentosa, to help preserve or restore vision in patients suffering from severe degenerative retinal diseases. GenSight Biologics' lead product candidate, GS010, is in Phase III trials in Leber's Hereditary Optic Neuropathy (LHON), a rare mitochondrial disease that leads to irreversible low vision and legal blindness in teens and young adults. Using its gene therapy-based approach, GenSight Biologics' product candidates are designed to be administered in a single treatment to each eye by intravitreal injection to offer patients a sustainable functional visual recovery.

Disclaimer

This document and the information contained herein do not constitute either an offer to sell or purchase, or the solicitation of an offer to sell or purchase, securities of GenSight Biologics S.A. (the "Company").

No communication and no information in respect of the offering by the Company of its shares may be distributed to the public in any jurisdiction where registration or approval is required. No steps have been taken or will be taken in any jurisdiction outside France where such steps would be required. The offering or subscription of shares may be subject to specific legal or regulatory restrictions in certain jurisdictions.

This announcement does not, and shall not, in any circumstances, constitute a public offering nor an invitation to the public in connection with any offer. The distribution of this document may be restricted by law in certain jurisdictions. Persons into whose possession this document comes are required to inform themselves about and to observe any such restrictions.

This announcement is an advertisement and not a prospectus within the meaning of the Prospectus Directive (as defined below), as implemented in each member State of the European Economic Area.

With respect to the Member States of the European Economic Area (including France) ("Member States"), no action has been undertaken or will be undertaken to make an offer to the public of the securities referred to herein requiring a publication of a prospectus in any Member State. As a result, the securities of the Company may not and will not be offered in any Member State except in accordance with the exemptions set forth in Article 3 of the Prospectus Directive.

For the purposes of the provision above, the expression "offer to the public" in relation to any shares of the Company in any Member State means the communication in any form and by any means of sufficient information on the terms of the offer and any securities to be offered so as to enable an investor to decide to purchase any securities, as the same may be varied in that Member State. The expression "Prospectus Directive" means Directive 2003/71/EC (as amended, including by Directive 2010/73/EU), and includes any relevant implementing measure in the Member State.

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